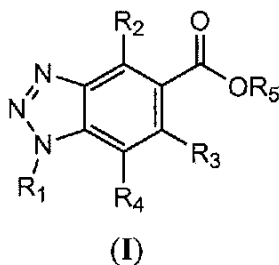


Amendments to the Claims:

Please cancel claim 21 and add new claim 29. Please amend claims 3-12, 21, 26, and 27 as shown herein. This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

1. (Previously presented) A compound of Formula (I):



wherein:

R<sub>1</sub> is C<sub>1-8</sub> alkyl, C<sub>3-6</sub> cycloalkyl or C<sub>1-6</sub> haloalkyl, wherein the C<sub>1-8</sub> alkyl, C<sub>3-6</sub> cycloalkyl and C<sub>1-6</sub> haloalkyl groups are optionally substituted with 1, 2, 3 or 4 substituents selected from the group consisting of C<sub>1-6</sub> acyl, C<sub>1-6</sub> acyloxy, C<sub>2-6</sub> alkenyl, C<sub>1-6</sub> alkoxy, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkylcarboxamido, C<sub>2-6</sub> alkynyl, C<sub>1-6</sub> alkylsulfinyl, C<sub>1-6</sub> alkylsulfonyl, C<sub>1-6</sub> alkylthio, C<sub>1-6</sub> alkylureyl, amino, C<sub>1-6</sub> alkylamino, aryl, substituted aryl, C<sub>1-6</sub> dialkylamino, carbo C<sub>1-6</sub> alkoxy, carboxy, cyano, C<sub>3-6</sub> cycloalkyl, C<sub>1-6</sub> dialkylcarboxamido, halogen, C<sub>1-6</sub> haloalkoxy, C<sub>1-6</sub> haloalkyl, C<sub>1-6</sub> haloalkylsulfinyl, C<sub>1-6</sub> haloalkylsulfonyl, C<sub>1-6</sub> haloalkylthio, heteroaryl, heterocyclyl, hydroxyl, nitro and thiol;

R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are each independently selected from the group consisting of H, C<sub>1-6</sub> acyl, C<sub>1-6</sub> acyloxy, C<sub>2-6</sub> alkenyl, C<sub>1-6</sub> alkoxy, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkylcarboxamido, C<sub>2-6</sub> alkynyl, C<sub>1-6</sub> alkylsulfinyl, C<sub>1-6</sub> alkylsulfonyl, C<sub>1-6</sub> alkylthio, C<sub>1-6</sub> alkylureyl, amino, C<sub>1-6</sub> alkylamino, C<sub>1-6</sub> dialkylamino, carbo C<sub>1-6</sub> alkoxy, carboxy, cyano, C<sub>3-6</sub> cycloalkyl, C<sub>1-6</sub> dialkylcarboxamido, halogen, C<sub>1-6</sub> haloalkoxy, C<sub>1-6</sub> haloalkyl, C<sub>1-6</sub> haloalkylsulfinyl, C<sub>1-6</sub> haloalkylsulfonyl, C<sub>1-6</sub> haloalkylthio, hydroxyl, nitro and thiol; and

R<sub>5</sub> is H or C<sub>1-6</sub> alkyl; or

a pharmaceutically acceptable salt, solvate or hydrate thereof;

provided that:

- a) when  $R_5$  is ethyl, and  $R_2$ ,  $R_3$  and  $R_4$  are H then  $R_1$  is not methyl or triphenylmethyl;
- b) when  $R_5$  is n-pentyl, and  $R_2$ ,  $R_3$  and  $R_4$  are H then  $R_1$  is not n-butyl;
- c) when  $R_5$  is methyl, and  $R_2$ ,  $R_3$  and  $R_4$  are H then  $R_1$  is not pyrrolidin-1-ylmethyl, 3-tert-butyl-2-hydroxy-5-methyl-benzyl, methyl, or dimethylaminomethyl;
- d) when  $R_5$  is methyl,  $R_2$  is carbomethoxy and  $R_3$  and  $R_4$  are both H then  $R_1$  is not methyl;
- e) when  $R_2$ ,  $R_3$ ,  $R_4$  and  $R_5$  are all H then  $R_1$  is not 2-amino-2-carboxy-ethyl, pyrrolidin-1-ylmethyl, isopropyl, methyl, benzyl, n-butyl, or carboxymethyl; and
- f) when  $R_2$ ,  $R_4$ , and  $R_5$  are all H and  $R_3$  is methoxy then  $R_1$  is not methyl.

2. (Previously presented) A compound according to claim 1 wherein:

$R_1$  is  $C_{3-6}$  cycloalkyl or  $C_{1-6}$  haloalkyl, where each  $C_{3-6}$  cycloalkyl or  $C_{1-6}$  haloalkyl group is optionally substituted with 1, 2, 3, or 4 substituents selected from the group consisting of  $C_{1-6}$  acyl,  $C_{1-6}$  acyloxy,  $C_{2-6}$  alkenyl,  $C_{1-6}$  alkoxy,  $C_{1-6}$  alkyl,  $C_{1-6}$  alkylcarboxamido,  $C_{2-6}$  alkynyl,  $C_{1-6}$  alkylsulfinyl,  $C_{1-6}$  alkylsulfonyl,  $C_{1-6}$  alkylthio,  $C_{1-6}$  alkylureyl, amino,  $C_{1-6}$  alkylamino,  $C_{1-6}$  dialkylamino, carbo  $C_{1-6}$  alkoxy, carboxy, cyano,  $C_{3-6}$  cycloalkyl,  $C_{1-6}$  dialkylcarboxamido, halogen,  $C_{1-6}$  haloalkoxy,  $C_{1-6}$  haloalkyl,  $C_{1-6}$  haloalkylsulfinyl,  $C_{1-6}$  haloalkylsulfonyl,  $C_{1-6}$  haloalkylthio, hydroxyl, nitro and thiol;

$R_2$ ,  $R_3$  and  $R_4$  are each independently selected from the group consisting of H,  $C_{1-6}$  acyl,  $C_{1-6}$  acyloxy,  $C_{2-6}$  alkenyl,  $C_{1-6}$  alkoxy,  $C_{1-6}$  alkyl,  $C_{1-6}$  alkylcarboxamido,  $C_{2-6}$  alkynyl,  $C_{1-6}$  alkylsulfinyl,  $C_{1-6}$  alkylsulfonyl,  $C_{1-6}$  alkylthio,  $C_{1-6}$  alkylureyl, amino,  $C_{1-6}$  alkylamino,  $C_{1-6}$  dialkylamino, carbo  $C_{1-6}$  alkoxy, carboxy, cyano,  $C_{3-6}$  cycloalkyl,  $C_{1-6}$  dialkylcarboxamido, halogen,  $C_{1-6}$  haloalkoxy,  $C_{1-6}$  haloalkyl,  $C_{1-6}$  haloalkylsulfinyl,  $C_{1-6}$  haloalkylsulfonyl,  $C_{1-6}$  haloalkylthio, hydroxyl, nitro or thiol; and

$R_5$  is H or  $C_{1-6}$  alkyl; or a pharmaceutically acceptable salt, solvate or hydrate thereof.

3. (Currently amended) The compound according to claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof, wherein  $R_5$  is  $C_{1-6}$  alkyl.

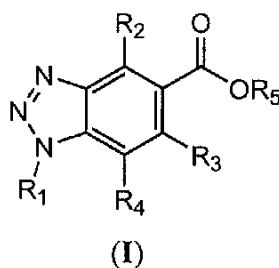
4. (Currently amended) The compound according to claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof, wherein R<sub>5</sub> is H.
5. (Currently amended) The compound according to claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof, wherein R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are each independently H or halogen.
6. (Currently amended) The compound according to claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof, wherein R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are each independently H or F.
7. (Currently amended) The compound according to claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof, wherein R<sub>1</sub> is C<sub>1-8</sub> alkyl optionally substituted with 1, 2, 3 or 4 substituents selected from the group consisting of C<sub>2-6</sub> alkenyl, C<sub>1-6</sub> alkoxy, C<sub>2-6</sub> alkynyl, C<sub>1-6</sub> alkylsulfinyl, C<sub>1-6</sub> alkylsulfonyl, C<sub>1-6</sub> alkylthio, aryl, substituted aryl, C<sub>3-6</sub> cycloalkyl, halogen, C<sub>1-6</sub> haloalkoxy, C<sub>1-6</sub> haloalkylsulfinyl, C<sub>1-6</sub> haloalkylsulfonyl, C<sub>1-6</sub> haloalkylthio, heteroaryl, heterocyclyl, and hydroxyl.
8. (Currently amended) The compound according to claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof, wherein R<sub>1</sub> is selected from the group consisting of 2-butyl, 3-pentyl, 1-propyl, t-butyl, 1-butyl, 4-Methyl-pentyl, 3-methyl-butyl, 1,3-dimethyl-butyl, 3,3-dimethyl-butyl, 1-heptyl, ethyl, 2,2-dimethyl-propyl, and 1-pentyl.
9. (Currently amended) The compound according to claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof, wherein R<sub>1</sub> is selected from the group consisting of 3-methoxy-benzyl, 4-methoxy-benzyl, 4-methoxy-phenyl ethyl, 3-methoxy-phenyl ethyl, 3,5-difluorobenzyl, and benzhydryl.
10. (Currently amended) The compound according to claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof, wherein R<sub>1</sub> is selected from the group consisting of 3-isopropoxypropyl, tetrahydro-furan-2-ylmethyl, 2-methoxy-ethyl, 2-ethylsulfonyl-ethyl, 3-

hydroxy-propyl, allyl, cyclopropylmethyl, but-2-ynyl, 2-methoxy-1-methyl-ethyl, 2-hydroxy-1-hydroxymethyl-ethyl, 2-ethoxy-ethyl, and 1,2-dimethyl-propyl.

11. (Currently amended) The compound according to claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof, wherein R<sub>1</sub> is selected from the group consisting of cyclopentyl, cyclohexyl, cyclopropyl, and cyclobutyl.
12. (Currently amended) The compound according to claim 1 selected from the group consisting of:
- 1-Cyclopentyl-1H-benzotriazole-5-carboxylic acid;
  - 1-(2'-Butyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-(3'-Pentyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-Cyclohexyl-1H-benzotriazole-5-carboxylic acid
  - 1-Propyl-1H-benzotriazole-5-carboxylic acid;
  - 1-Cyclopropyl-1H-benzotriazole-5-carboxylic acid;
  - 1-(3'-Isopropoxy-propyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-(Tetrahydro-furan-2'-ylmethyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-Cyclobutyl-1H-benzotriazole-5-carboxylic acid;
  - 1-(2-Methoxy-ethyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-(3'Methoxybenzyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-(4'Methoxybenzyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-[2'-(4''-Methoxy-phenyl)-ethylamino]-1H-benzotriazole-5-carboxylic acid;
  - 1-[2'-(3''-Methoxy-phenyl)-ethylamino]-1H-benzotriazole-5-carboxylic acid;
  - 1-(3',5'-Difluorobenzyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-(2-Ethylsulfanyl-ethyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-t-Butyl-1H-benzotriazole-5-carboxylic acid;
  - 1-(3'-Hydroxy-propyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-(1',3'-Dimethyl-butyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-(3',3'-Dimethyl-butyl)-1H-benzotriazole-5-carboxylic acid;
  - 1-Heptyl-1H-benzotriazole-5-carboxylic acid;
  - 1-(2'-Methoxy-1'-methyl-ethyl)-1H-benzotriazole-5-carboxylic acid;

1-(2'-Hydroxy-1'-hydroxymethyl-ethyl)-1H-benzotriazole-5-carboxylic acid;  
1-Ethyl-1H-benzotriazole-5-carboxylic acid;  
1-Pentyl-1H-benzotriazole-5-carboxylic acid;  
1-(2',2'-Dimethyl-propyl)-1H-benzotriazole-5-carboxylic acid;  
1-(2'-Ethoxy-ethyl)-1H-benzotriazole-5-carboxylic acid;  
1-(1',2'-Dimethyl-propyl)-1H-benzotriazole-5-carboxylic acid;  
1-Benzhydryl-1H-benzotriazole-5-carboxylic acid;  
1-Allyl-1H-benzotriazole-5-carboxylic acid;  
~~1-Butyl-1H-benzotriazole-5-carboxylic acid;~~  
1-(Cyclopropylmethyl)-1H-benzotriazole-5-carboxylic acid;  
1-(But-2-ynyl)-1H-benzotriazole-5-carboxylic acid;  
1-(4'-Methyl-pentyl)-1H-benzotriazole-5-carboxylic acid; and  
1-(3'-Methyl-butyl)-1H-benzotriazole-5-carboxylic acid; or  
a pharmaceutically acceptable salt, solvate or hydrate thereof.

13. (Previously presented) A pharmaceutical composition comprising a compound according to Formula (I):



wherein:

R<sub>1</sub> is H, C<sub>1-8</sub> alkyl, C<sub>3-6</sub> cycloalkyl or C<sub>1-6</sub> haloalkyl, wherein each C<sub>1-6</sub> alkyl, C<sub>3-6</sub> cycloalkyl or C<sub>1-6</sub> haloalkyl group is optionally substituted with 1, 2, 3, or 4 substituents selected from the group consisting of C<sub>1-6</sub> acyl, C<sub>1-6</sub> acyloxy, C<sub>2-6</sub> alkenyl, C<sub>1-6</sub> alkoxy, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkylcarboxamido, C<sub>2-6</sub> alkynyl, C<sub>1-6</sub> alkylsulfinyl, C<sub>1-6</sub> alkylsulfonyl, C<sub>1-6</sub> alkylthio, C<sub>1-6</sub> alkylureyl, amino, C<sub>1-6</sub> alkylamino, C<sub>1-6</sub> dialkylamino, carbo C<sub>1-6</sub> alkoxy, carboxy, cyano, C<sub>3-6</sub> cycloalkyl, C<sub>1-6</sub> dialkylcarboxamido, halogen, C<sub>1-6</sub> haloalkoxy, C<sub>1-6</sub> haloalkyl, C<sub>1-6</sub> haloalkylsulfinyl, C<sub>1-6</sub> haloalkylsulfonyl, C<sub>1-6</sub> haloalkylthio, hydroxyl, nitro or thiol;

R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are each independently selected from the group consisting of H, C<sub>1-6</sub> acyl, C<sub>1-6</sub> acyloxy, C<sub>2-6</sub> alkenyl, C<sub>1-6</sub> alkoxy, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkylcarboxamido, C<sub>2-6</sub> alkynyl, C<sub>1-6</sub> alkylsulfinyl, C<sub>1-6</sub> alkylsulfonyl, C<sub>1-6</sub> alkylthio, C<sub>1-6</sub> alkylureyl, amino, C<sub>1-6</sub> alkylamino, C<sub>1-6</sub> dialkylamino, carbo C<sub>1-6</sub> alkoxy, carboxy, cyano, C<sub>3-6</sub> cycloalkyl, C<sub>1-6</sub> dialkylcarboxamido, halogen, C<sub>1-6</sub> haloalkoxy, C<sub>1-6</sub> haloalkyl, C<sub>1-6</sub> haloalkylsulfinyl, C<sub>1-6</sub> haloalkylsulfonyl, C<sub>1-6</sub> haloalkylthio, hydroxyl, nitro or thiol; and

R<sub>5</sub> is H or C<sub>1-6</sub> alkyl; or

a pharmaceutically acceptable salt, solvate or hydrate thereof, in combination with a pharmaceutically acceptable carrier.

14. (Original) A pharmaceutical composition according to claim 13 further comprising an agent selected from the group consisting of  $\alpha$ -glucosidase inhibitor, aldose reductase inhibitor, biguanide, HMG-CoA reductase inhibitor, squalene synthesis inhibitor, fibrate, LDL catabolism enhancer, angiotensin converting enzyme inhibitor, insulin secretion enhancer and thiazolidinedione.

15-19. (Canceled)

20. (Currently amended) A method of treatment of a metabolic-related disorder selected from the group consisting of dyslipidemia, atherosclerosis, coronary heart disease, insulin resistance and type 2 diabetes comprising administering to an individual in need of such treatment a therapeutically effective amount of a pharmaceutical composition according to claim 13.

21-25. (Canceled)

26. (Currently amended) A method of treatment of a metabolic-related disorder selected from the group consisting of dyslipidemia, atherosclerosis, coronary heart disease, insulin resistance and type 2 diabetes comprising administering to an individual in need of such treatment a therapeutically effective amount of a compound according to claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof.

27. (Currently Amended) A method according to claim 26 wherein said metabolic-related disorder is selected from the group consisting of dyslipidemia, atherosclerosis, coronary heart disease, insulin resistance and type 2 diabetes.

28. (Previously presented) The pharmaceutical composition according to claim 13 wherein said compound is selected from the group consisting of:

- 1-Isopropyl-1H-benzotriazole-5-carboxylic acid;
- 1-Cyclopentyl-1H-benzotriazole-5-carboxylic acid;
- 1-(2'-Butyl)-1H-benzotriazole-5-carboxylic acid;
- 1-(3'-Pentyl)-1H-benzotriazole-5-carboxylic acid;
- 1-Benzyl-1H-benzotriazole-5-carboxylic acid;
- 1-Cyclohexyl-1H-benzotriazole-5-carboxylic acid
- 1-Propyl-1H-benzotriazole-5-carboxylic acid;
- 1-Cyclopropyl-1H-benzotriazole-5-carboxylic acid;
- 1-(3'-Isopropoxy-propyl)-1H-benzotriazole-5-carboxylic acid;
- 1-(Tetrahydro-furan-2'-ylmethyl)-1H-benzotriazole-5-carboxylic acid;
- 1-Cyclobutyl-1H-benzotriazole-5-carboxylic acid;
- 1-(2-Methoxy-ethyl)-1H-benzotriazole-5-carboxylic acid;
- 1-(3'Methoxybenzyl)-1H-benzotriazole-5-carboxylic acid;
- 1-(4'Methoxybenzyl)-1H-benzotriazole-5-carboxylic acid;
- 1-[2'-(4''-Methoxy-phenyl)-ethylamino]-1H-benzotriazole-5-carboxylic acid;
- 1-[2'-(3''-Methoxy-phenyl)-ethylamino]-1H-benzotriazole-5-carboxylic acid;
- 1-(3',5'-Difluorobenzyl)-1H-benzotriazole-5-carboxylic acid;
- 1-(2-Ethylsulfanyl-ethyl)-1H-benzotriazole-5-carboxylic acid;
- 1-t-Butyl-1H-benzotriazole-5-carboxylic acid;
- 1-(3'-Hydroxy-propyl)-1H-benzotriazole-5-carboxylic acid;
- 1-(1',3'-Dimethyl-butyl)-1H-benzotriazole-5-carboxylic acid;
- 1-(3',3'-Dimethyl-butyl)-1H-benzotriazole-5-carboxylic acid;
- 1-Heptyl-1H-benzotriazole-5-carboxylic acid;
- 1-(2'-Methoxy-1'-methyl-ethyl)-1H-benzotriazole-5-carboxylic acid;
- 1-(2'-Hydroxy-1'-hydroxymethyl-ethyl)-1H-benzotriazole-5-carboxylic acid;

1-Ethyl-1H-benzotriazole-5-carboxylic acid;  
1-Pentyl-1H-benzotriazole-5-carboxylic acid;  
1-(2',2'-Dimethyl-propyl)-1H-benzotriazole-5-carboxylic acid;  
1-(2'-Ethoxy-ethyl)-1H-benzotriazole-5-carboxylic acid;  
1-(1',2'-Dimethyl-propyl)-1H-benzotriazole-5-carboxylic acid;  
1-Benzhydryl-1H-benzotriazole-5-carboxylic acid;  
1-Allyl-1H-benzotriazole-5-carboxylic acid;  
1-Butyl-1H-benzotriazole-5-carboxylic acid;  
1-(Cyclopropylmethyl)-1H-benzotriazole-5-carboxylic acid;  
1-(But-2-ynyl)-1H-benzotriazole-5-carboxylic acid;  
1-(4'-Methyl-pentyl)-1H-benzotriazole-5-carboxylic acid; and  
1-(3'-Methyl-butyl)-1H-benzotriazole-5-carboxylic acid; or  
a pharmaceutically acceptable salt, solvate or hydrate thereof.

29. (New) A method according to claim 26 wherein said metabolic-related disorder is atherosclerosis.